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	Application Number		10587467	
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INFORMATION DISCLOSURE	First Named Inventor	tor Maruoka		
STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)	Art Unit		1044 1624	
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/B.C./	1	BELLIER, et al., Synthesis and Biological Properties of New Constrained CCK-B Antagonists: Discrimination of Two Affinity States of the CCK-B Receptor on Transfected CHO Cells, J. Med. Chem., Vol. 40, No. 24, pp.3947-3956, 1997	
/B.C./	2	SHIOIRI, et al., Asymmetric Phase Transfer Catalysis, Stimulating Concepts in Chemistry, pp.123-143, 2000, Wiley-VCH, Weinheim	
/B.C./	3	OOI, et al., Practical Catalytic Enantioselective Synthesis of a,a-Dialkyl-a-Amino Acids by Chiral Phase-Transfer Catalysis, J. Am. Chem. Soc., Vol. 122, pp. 5228-5229, 2000	
/B.C./	4	SEKI, et al., A Practical Synthesis of C2-Symmetric Chiral Binaphthyl Ketone Catalyst, Synthesis, No. 12, pp. 1677-1680, 2000	
/B.C./	5	OOI, et al., New, Improved Procedure for the Synthesis of Structurally Diverse N-Spiro C2-Symmetric Chiral Quaternary Ammonium Bromides, J. Org. Chem., Vol. 68, pp. 4576-4578, 2003	
/B.C./	6	OOI, et al., Design of N-Spiro C2-Symmetric Chiral Quaternary Ammonium Bromides as Novel Chiral Phase-Transfer Catalysts: Synthesis and Application to Practical Asymmetric Synthesis of a-Amino Acids, J. Am. Chem. Soc., Vol. 125, No. 17, pp. 5139-5151, 2003	
/B.C./	7	OOI, et al., Molecular Design of a C2-Symmetric Chiral Phase-Transfer Catalyst for Practical Asymmetric Synthesis of a-Amino Acids, J. Am. Chem. Soc., Vol. 121, No. 27, pp. 6519-6520, 1999	
/B.C./	8	STARA, et al., Nucleophilic Cleavage of 4,5-Dihydro-3H-dinaphth[2,1-c:1',2'-e]azepinium Quaternary Salts. A Convenient Approach to New Axially Dissymmetric and Axially Asymmetric Ligands, J. Org. Chem., Vol. 57, No. 25, pp. 6966-6969, 1992	
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/B.C./	12	2 IKUNAKA, et al., A Scalable Synthesis of (R)-3,5-Dihydro-4H-dinaphth[2,1-c:1'2'-e]azepine, Organic Process Research & Development, Vol. 7, No. 5, pp. 644-648, 2003					
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